A new approach in the assessment of motor activity in Parkinson’s disease

J J van Hilten, H A M Middelkoop, G A Kerkhof, R A C Roos

Abstract
Motor activity was recorded continuously with a wrist-worn activity monitor for almost six days in nine patients with different predominant manifestations of Parkinson's disease and 10 normal subjects. The indices of diurnal motor activity decreased with the progressive worsening of hypokinesia and rigidity. With this monitor and a simple diary it was possible to determine the contribution of a moderate resting tremor and choreiform dyskinesia to the motor activity level, and to monitor their variability.

In the clinical assessment of motor symptoms in Parkinson's disease (PD), both simple motor tests and complex neurophysiological recordings are used. These tests are stressful for the patient, time-consuming, costly, and assess only the momentary extent of the different motor symptoms, which renders them particularly susceptible to the influence of stress. The reliability of functional motor assessment in PD is therefore seriously hampered by fluctuations in severity of symptoms. In addition, diaries may be used, in which the patient records the duration and severity of motor symptoms. However, these diaries are cumbersome and usually not completed.

There is a need for an objective continuous, measurement procedure of motor activity which does not interfere with daily activities. Various wrist-worn activity monitors have been introduced and validated in sleep, epidemiological and behavioural studies. The activity monitor promises to be an excellent tool for the type of recording needed. However, it cannot discriminate motor activity caused by volitional movements, tremor or dyskinesia. Furthermore, with activity monitoring diurnal immobility periods due to hypokinesia are indistinguishable from naps. Thus depending on the monitor design, that is, sensitivity to a certain frequency range, this method relies partly on the traditional assessment of Parkinsonian symptoms and the keeping of a simple diary in which tremor and/or dyskinesia as well as daytime naps and sleep are indicated. We describe our preliminary results with the activity monitor.

Patients and methods
Nine patients (six males and three females) with idiopathic Parkinson's disease [age 44–80, (mean 68) years] were compared with ten healthy subjects [five males and five females, age 55–80 (mean 70) years]. Five PD patients (patients 1–5) are characterised by hypokinesia and rigidity and show a stable levodopa response. Two PD patients (patients 6 and 7) also had a severe resting tremor. Two PD patients (patients 8 and 9) with young-onset had severe on-off fluctuations with choreiform dyskinesia. Clinical data are presented in table 1. Supplementary to levodopa patients received other anti-Parkinsonian medications, including trihexyphenidyl hydrochloride (patient 5), bromocriptine (patients 5, 6), and lisuride (patients 8, 9). Patients with a Mini-Mental State Examination score below 24 were excluded. All patients were clinically assessed by the Hoehn and Yahr and Unified Parkinson's Disease Rating Scale (UPDRS). The score of the motor examination (UPDRS) and presence and severity of a resting tremor on the nondominant side are noted separately (table 1).

In all subjects motor activity was recorded by means of a monitor, worn on the nondominant wrist from Monday 7 pm until Sunday 11 am. Subjects were asked to maintain their habitual 24 hour pattern of activities, and to remove the monitor only when taking a bath. During the recording period all subjects kept a log in which they recorded the time they switched off the light to

Table 1  Clinical characteristics of nine patients with Parkinson’s disease

<table>
<thead>
<tr>
<th>Patient number</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
<th>7</th>
<th>8</th>
<th>9</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age of onset (years)</td>
<td>64</td>
<td>76</td>
<td>67</td>
<td>68</td>
<td>61</td>
<td>70</td>
<td>52</td>
<td>37</td>
<td>49</td>
</tr>
<tr>
<td>Duration of illness (years)</td>
<td>0.8</td>
<td>2</td>
<td>5</td>
<td>3</td>
<td>13</td>
<td>10</td>
<td>13</td>
<td>7</td>
<td>15</td>
</tr>
<tr>
<td>Levodopa treatment (years)</td>
<td>0</td>
<td>0.5</td>
<td>3</td>
<td>0</td>
<td>10</td>
<td>0.4</td>
<td>0.4</td>
<td>1.1</td>
<td>1.75</td>
</tr>
<tr>
<td>Average dosage levodopa (g/day)</td>
<td>0.25</td>
<td>0.3</td>
<td>0.4</td>
<td>0.3</td>
<td>0.4</td>
<td>0.4</td>
<td>0.4</td>
<td>0.4</td>
<td>0.4</td>
</tr>
<tr>
<td>Hoehn and Yahr (off)</td>
<td>2</td>
<td>3</td>
<td>3</td>
<td>2</td>
<td>4</td>
<td>3</td>
<td>4</td>
<td>4</td>
<td>4</td>
</tr>
<tr>
<td>UPDRS (on)</td>
<td>3</td>
<td>35</td>
<td>34</td>
<td>43</td>
<td>53</td>
<td>61</td>
<td>67</td>
<td>74</td>
<td></td>
</tr>
<tr>
<td>UPDRS motor examination score</td>
<td>15</td>
<td>19</td>
<td>19</td>
<td>28</td>
<td>33</td>
<td>32</td>
<td>28</td>
<td>36</td>
<td>46</td>
</tr>
<tr>
<td>Tremor sub-score</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>3</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

UPDRS = the Unified Parkinson's Disease Rating Scale
go to sleep, the time of final awakening, any naps, and the time the monitor was removed when taking a bath.

The characteristics of the monitor (Gaehwiler Electronic, CH-8634 Hombrechtikon) used in this study have been described previously. The device counts the occurrence of supra-threshold motor activity (accelerations > 0.1 g with filtering of the analog sensor signal by a bandpass filter of 0.25–3 Hz) over 30-second-epochs, and stores the resulting sum as a 1-byte value in a 32 kbyte solid state memory. After completion of the recording, the data were read out on computer and analysed further.

For each recording period the following parameters of motor activity were calculated ("diurnal" is defined as the time from final awakening to lights-out): 1) The diurnal activity level (DAL), expressed as the mean number of counts per 30 seconds calculated over all days; 2) The diurnal movement index (DMI), calculated as the mean over all days. The diurnal movement index of each day is the number of 30-second-epochs with any movement (activity count > 0) expressed as percentage of 30-second-epochs comprising the diurnal period. In contrast with the DAL, the DMI reflects the degree of clustering of activity and inactivity; that is, a relatively small value indicates that motor activity only occurs during a restricted part(s) of the day; 3) The variability of the DAL and DMI during the recording period, expressed as the coefficient of variation (CV); CV-DAL and CV-DMI, respectively.

Because the graphics software (Harvard graphics) allows for only 240 X-axis values, the graphs show activity counts per six minutes which are obtained by adding the counts of 12 successive 30-second-epochs.

**Results**

**Normal subjects**

A 24 hour plot of the overall mean (SD) values of the monitor counts for the normal subjects is presented in fig 1A. Three levels of diurnal activity may be observed: 1) the night (0:30 am–8 am), with the lowest level of motor activity; 2) the day (8 am–6.30 pm), which shows a gradual increase of motor activity between 8–9 am and a more stable level with the highest values between 10 am and 6.30 pm; 3) the evening (6.30 pm–0.30 am), which shows an intermediate level of motor activity with a gradual decline. Table 2 gives the values of the various parameters of motor activity.

### Table 2: Diurnal motor activity characteristics of nine patients with Parkinson's disease and mean values of 10 normal subjects. Motor activity expressed as activity counts/30 seconds

<table>
<thead>
<tr>
<th>Normal subjects</th>
<th>Parkinson patients with Hypokinesia and rigidity</th>
<th>Resting tremor</th>
<th>Response fluctuation</th>
</tr>
</thead>
<tbody>
<tr>
<td>DAL 75</td>
<td>55 38 28 10 3</td>
<td>77 125</td>
<td>50 105</td>
</tr>
<tr>
<td>CV-DAL (%) 17</td>
<td>44 38 29 27 79</td>
<td>19 31 22</td>
<td>22 9</td>
</tr>
<tr>
<td>DMI (%) 85</td>
<td>66 67 59 49 15</td>
<td>67 75 52</td>
<td>62 62</td>
</tr>
<tr>
<td>CV-DMI (%) 7</td>
<td>18 11 15 16 79</td>
<td>8 15 15</td>
<td>27 27</td>
</tr>
</tbody>
</table>

DAL = diurnal activity level, DMI = diurnal movement index, CV = coefficient of variation.
increased clustering of motor activity. The early afternoon dip in the curve is caused by a daily nap.

The graph of patient 1 (fig 1B) shows a rapid decline of motor activity in the course of the day and illustrates that conspicuous diurnal fluctuations in motor activity can be detected in untreated patients with early PD.

The graph of patient 9 (fig 3) clearly shows the regular switching of episodes with low and high motor activity. This patient has on-off fluctuations with choreiform dyskinesias, at a treatment regime of orally administered levodopa every four hours. Because of the severe choreiform dyskinesias the diurnal activity level (table 2) of this patient is tremendously elevated. However, as in the patients with tremor the diurnal movement index was lower, again indicating the increased clustering of motor activity.

Discussion

The advantages of the monitor largely derive from its potential to record unrestrained motor activity for several days continuously, not influenced by stress such as visiting the doctor. Our results show that each of the features of PD may contribute to the motor activity status as measured by the monitor. With the progression of hypokinesia and rigidity, the diurnal indices of motor activity decrease. On the other hand, the effects of resting tremor and choreiform dyskinesias are shown by the combined presence of a normal or elevated diurnal activity level associated with a lower diurnal movement index, indicating a relative clustering of motor activity.

In normal subjects voluntary movements do not occur at a rate greater than 200/minute (3-3 Hz).22 Because of hypokinesia and rigidity this rate will obviously be slower in patients with PD. However, in Parkinsonian patients with a predominant resting tremor, a monitor design which is specifically responsive to the frequency characteristics of the resting tremor (4-6 Hz) would be more desirable. In this study we were interested in whether the different motor features of PD could be monitored with the same monitor design during a continuous recording. Our results show that with the monitor design used it is possible to determine the extent to which a resting tremor which is clinically judged as moderate in amplitude and present most of the time, contributes to the motor activity level, and of its variability.

Because of the variability of symptom intensity, the assessment of the effectiveness of therapeutic interventions in PD is difficult. This applies to every stage of PD but particularly in the case of response fluctuations. The strength of this method is that the time of day effects of the on-off fluctuations can be studied, and that the amount of "off-time" can be quantified. However, the monitor data are useful if interpreted with the information obtained by a diary. Without a diary it is impossible to distinguish between the various sources of decreased (naps or severe hypo-

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*Figure 2* (patient 7). The mean 24 hour motor activity profile (over a period of almost six days) of a patient with Parkinson's disease with tremor as a predominant manifestation. The solid line represents the grand mean of motor activity of the 10 normal subjects.

*Figure 3* (patient 9). The mean 24 hour motor activity profile (over a period of almost six days) of a patient with Parkinson's disease suffering from on-off fluctuations with choreiform dyskinesias. Treatment regime of levodopa administered orally every four hours. The solid line represents the grand mean of motor activity of the 10 normal subjects.
A new approach in the assessment of motor activity in Parkinson's disease

A new approach in the assessment of motor activity in Parkinson's disease (kinesia) or elevated (tremor or chorea) motor activity.

In conclusion, despite some disadvantages the monitor is a welcome adjunct to the more subjective assessments of PD because of its simplicity and practicality in all settings. The monitor may prove to be of great value for studies of the disease course and response to new drug regimes.

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